This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. - 11. (Cancelled)

- 12. (New) A synthesized peptide comprising one or more sequences selected from the group consisting of SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, SEQ ID NO:8 and SEQ ID NO:11.
- 13. (New) An osteogenetic accelerator comprising the peptide set forth in claim 12, or a pharmacologically acceptable salt thereof, attached to a biocompatible carrier.
- 14. (New) An osteogenetic accelerator comprising the peptide of claim 12, or a pharmacologically acceptable salt thereof, mixed with, dissolved in, or suspended in aqueous solvent.
 - 15. (New) A synthesized peptide comprising the sequence SEQ ID NO:11.
- 16. (New) An osteogenetic accelerator comprising the peptide set forth in claim 15, or a pharmacologically acceptable salt thereof, attached to a biocompatible carrier.
- 17. (New) The osteogenetic accelerator according to claim 16, wherein the carrier is selected from a group consisting of a ceramic, an artificial bone, a covalently cross-linked gel of alginate, and a gel of collagen, hyaluronic acid,

calcium sulfate, polylactic acid, polyglycolic acid, hydroxyapatite, tricalcium phosphate, starch, chitin/chitosan, agarose, or dextran.

- 18 (New) The osteogenetic accelerator according to claim 16 which contains 0.01 to 50 parts by weight of the peptide per 100 parts by weight of the carrier.
- 19. (New) An osteogenetic accelerator comprising the peptide of claim 15, or a pharmacologically acceptable salt thereof, mixed with, dissolved in, or suspended in aqueous solvent.
- 20. (New) The osteogenetic accelerator according to claim 19, wherein the aqueous solvent is physiological saline solution or a physiologically acceptable aqueous solution selected from a group consisting of mannitol, sucrose, lactose, maltose, glucose, and fructose.
- 21. (New) The osteogenetic accelerator according to claim 19 or 20, wherein the concentration of the peptide is 0.001% to 5% with respect to the aqueous solvent.
- 22. (New) The osteogenetic accelerator as set forth in claim 16 which is used for treating a bone fracture by inducing bone formation at the fracture site or for inhibiting a decrease in bone substance.
- 23. (New) The peptide of claim 15, wherein the peptide N-terminal is acetylated, or the peptide C-terminal is amidated, or both the N-terminal is acetylated and the C-terminal is amidated.

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24. (New) An osteogenetic accelerator comprising a physiologically acceptable salt of the peptide set forth in claim 15.

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